

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Administration for Children and Families****Head Start Bureau; Advisory Committee on Head Start Research and Evaluation; Meeting**

AGENCY: Administration for Children, Youth and Families, ACF, DHHS.

ACTION: Notice of meeting; Advisory Committee on Head Start Research and Evaluation.

SUMMARY: The 1998 Head Start Reauthorization (42 U.S.C. 9844(g); section 649(g)(1) of the Head Start Act, as amended) called on the Secretary of Health and Human Services to form an independent panel of experts (i.e., an Advisory Committee) to offer advice concerning research designs that would provide a national analysis of the impact of Head Start Programs. The July 26–27, 1999 meeting will be the last of three meetings of the Advisory Committee that will culminate in a report to the Secretary due September 30, 1999.

DATES: July 26, 1999, 9 a.m.–5 p.m. and July 27, 1999, 9 a.m.–5 p.m.

PLACE: Holiday Inn Hotel and Suites, 625 First Street, Alexandria, VA 22314.

SUPPLEMENTARY INFORMATION: This meeting is open to the public and is barrier free. Meeting records will also be open to the public and will be kept at the Switzer Building located at 330 “C” Street, SW, Washington, DC 20447. The Head Start Bureau also intends to make material related to this meeting available on the Head Start web site <http://www2.acf.dhhs.gov/programs/hshb>. An interpreter for the deaf and hearing impaired will be available upon advance request by calling Ellsworth Associates at 703/821–3090 (ext. 282).

FOR FURTHER INFORMATION CONTACT: Deborah Roderick Stark at 301/889–0430 for substantive information. ACF Office of Public Affairs at 202/401–9215 for press inquiries. Ellsworth Associates at 703/821–3090 (ext. 282) for logistical information.

Dated: July 1, 1999.

Patricia Montoya,

Commissioner, Administration on Children, Youth and Families.

[FR Doc. 99–17237 Filed 7–6–99; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Food and Drug Administration**

[Docket No. 97D–0268]

Guidance for Industry on Container Closure Systems for Packaging Human Drugs and Biologics; Chemistry, Manufacturing, and Controls Documentation; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a guidance for industry entitled “Container Closure Systems for Packaging Human Drugs and Biologics; Chemistry, Manufacturing, and Controls Documentation.” This guidance provides recommendations on the container closure systems information that applicants should provide to the Center for Drug Evaluation and Research (CDER) and the Center for Biologics Evaluation and Research (CBER) in support of new drug applications, abbreviated new drug applications, biologics license applications, and supplements to these applications.

DATES: Written comments may be submitted at any time.

ADDRESSES: Copies of this guidance for industry are available on the Internet at “<http://www.fda.gov/cder/guidance/index.htm>” or “<http://www.fda.gov/cber/guidelines.htm>”. Submit written requests for single copies of the guidance for industry to the Drug Information Branch (HFD–210), Center for Drug Evaluation and Research, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, or to the Office of Communication, Training and Manufacturers Assistance (HFM–40), Center for Biologics Evaluation and Research, Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852–1448. Send one self-addressed adhesive label to assist the office in processing your requests. Submit written comments on the draft guidance to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT:

W. Mike Adams, Center for Drug Evaluation and Research (HFD–180), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7310, or

John D. Finkbohner, Center for Biologics Evaluation and Research

(HFM–676), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852, 301827–3031.

SUPPLEMENTARY INFORMATION: FDA is announcing the availability of a guidance for industry entitled “Container Closure Systems for Packaging Human Drugs and Biologics; Chemistry, Manufacturing, and Controls Documentation.” This guidance provides recommendations on the container closure system information that applicants should provide to CDER or CBER for initial applications and supplements. In addition, the document provides guidance on qualification and quality control of packaging components used for particular dosage forms and routes of administration, including the following: Drug products for injection and ophthalmic drug products, liquid-based oral and topical drug products and topical delivery systems, solid oral dosage forms and powders for reconstitution, and other dosage forms. This guidance supersedes the agency’s “Guideline for Submitting Documentation for Packaging for Human Drugs and Biologics,” issued February 1987.

This Level 1 guidance is being issued consistent with FDA’s good guidance practice (62 FR 8961, February 27, 1997). In the **Federal Register** of July 15, 1997 (62 FR 37925), FDA announced the availability of a draft version of this guidance. The July 1997 document gave interested persons an opportunity to submit comments through September 15, 1997. On September 5, 1997, in response to requests from the public, the agency extended the comment period until November 14, 1997 (62 FR 46980). All comments received during the comment period have been carefully reviewed and incorporated in this revised guidance where appropriate. As a result of public input during the comment period, the final guidance is clearer and more concise than the draft version. The guidance represents the agency’s current thinking on submitting information in drug applications on container closure systems used in packaging human drugs and biologics. It does not create or confer any rights for or on any person and does not operate to bind FDA or the public. An alternative approach may be used if such approach satisfies the requirements of the applicable statute, regulations, or both.

Interested persons may, at any time, submit written comments on the guidance to the Dockets Management Branch (address above). Two copies of